



**Europäisches Patentamt**  
**European Patent Office**  
**Office européen des brevets**



⑪ Publication number:

**0 507 392 A2**

12

**EUROPEAN PATENT APPLICATION**

②<sup>1</sup> Application number: 92200885.9

⑤ Int. Cl.<sup>5</sup>: **G01R 33/48**, **G01R 33/56**

② Date of filing: 30.03.92

③ Priority: 02.04.91 EP 91200762

④3 Date of publication of application:  
07.10.92 Bulletin 92/41

ⓑ Designated Contracting States:  
DE FR GB NL

**(71) Applicant: N.V. Philips' Gloeilampenfabrieken  
Groenewoudseweg 1  
NL-5621 BA Eindhoven(NL)**

72 Inventor: **KcKinnon, Graeme**  
c/o INT. OCTROOIBUREAU B.V. Prof.  
Holstlaan 6  
NL-5656 AA Eindhoven(NL)

**Inventor: Fischer, Stefan**  
**c/o INT. OCTROOIBUREAU B.V. Prof.**  
**Holstlaan 6**  
**NL-5656 AA Eindhoven(NL)**  
**Inventor: Boesiger, Peter**  
**c/o INT. OCTROOIBUREAU B.V. Prof.**  
**Holstlaan 6**  
**NL-5656 AA Eindhoven(NL)**

74 Representative: **De Jongh, Cornelis**  
**Dominicus et al**  
**INTERNATIONAAL OCTROOIBUREAU B.V.**  
**Prof. Holstlaan 6**  
**NL-5656 AA Eindhoven(NL)**

54 Magnetic resonance imaging method and device for monitoring motion of a part of an object.

⑤7) Magnetic resonance imaging methods for myocardial tagging are known for assessment of myocardial motion. In such methods tagging pulse and gradient sequences (tsq) are applied to the heart of an object (4) for acquiring tagged magnetic resonance images of, for example, heart slices from diastole to systole, the sequences being triggered upon detection of the R-wave of an ECG. With these known sequences, such as the so-called SPAMM sequence, it is very difficult, if at all possible, to automatically detect the tagging grid and therefore to evaluate the motion quantitatively.

According to the present invention, a magnetic resonance imaging method and device (1) are disclosed which allow for automatic quantitative motion evaluation of a part of an object (4). The pulse and gradient sequences (tsq) are applied in pairs with spatially differing tagging patterns, images corresponding to the pairs being subtracted so as to form a tagged image. The motion is monitored by scanning of the tagging pattern in the tagged image.

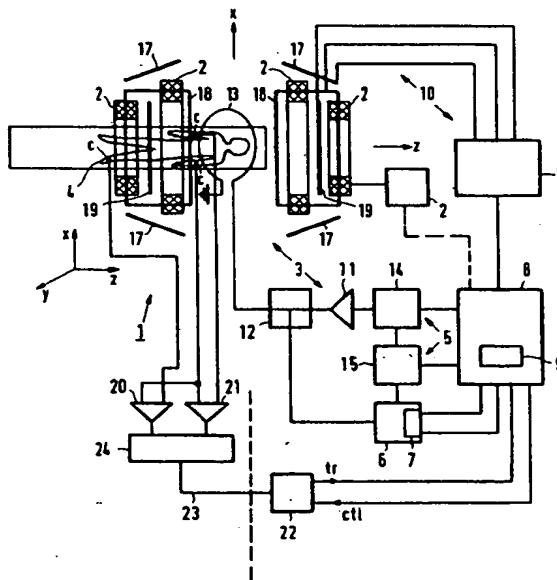


FIG. 1

**EP 0 507 392 A2**

The present invention relates to a magnetic resonance imaging method for monitoring motion of a part of an object to be imaged by application of pulse and gradient sequences to the object, which sequences comprise a tagging section for producing a tagging pattern and an imaging section for producing magnetic resonance signals in the part of the object, the resonance signals being used for forming at least one image of the object. Such tagging patterns can be used in connection with cardiac imaging as so-called myocardial tagging grids to evaluate heart motion in multi heart phase imaging, but can also be used for other purposes such as radiation therapy port planning of a liver. In the latter case the tagging is quasi static.

The present invention further relates to a magnetic resonance imaging device for monitoring motion of a part of an object, which imaging device comprises means for generating a stationary homogeneous magnetic field, means for radiating radio frequency pulses to the object which is placed in the stationary field, gradient generating means for generating magnetic gradient fields which are superimposed upon the stationary field, and processing means which are arranged for applying pulse and gradient sequences to the object under the control of control means, the sequences comprising a tagging section and an imaging section, and which imaging device further comprises receiving means and sampling means for receiving and sampling of magnetic resonance signals generated by the sequences in at least a part of the object, the processing means being arranged for processing sampled resonance signals into at least one image.

A method and device of this kind are known from the article, "MR Imaging of Motion with Spatial Modulation of Magnetization", L. Axel et al., Radiology, Volume 171, No. 3, pp. 841-845. For direct imaging of motion of the myocardium a multi heart phase imaging sequence is preceded by a pre-imaging so-called tagging sequence in order to obtain a line or grid reference pattern in the image. The tagging sequence is applied upon detection of the R-wave of an electrocardiogram. The multi heart phase imaging sequence produces magnetic resonance signals in a slice of the heart at different time intervals from the R-wave. Successive magnetic resonance images to be displayed visualize heart motion such as contraction and rotation, the tagging grid in the myocardium being displaced with respect to the chest wall tagging grid in successive heart images of the same slice. The tagging section of the imaging sequence comprises a radio frequency pulse to produce transverse magnetization, a magnetic field gradient to "wrap" the phase along the direction of the gradient and a further radio frequency pulse to mix the modulated

transverse magnetization with the longitudinal magnetization. The resulting images show periodic stripes due to the modulation. A second set of modulation bands, for example, in a direction orthogonal to the first, can be produced by following the further radio frequency pulse with a further magnetic field gradient in an appropriate direction and then with a still further radio frequency pulse. Different tagging sections have also been described, e.g. for obtaining a radial tagging pattern, in "A Rapid Starburst Pulse Sequence for Cardiac Tagging", E. McVeigh et al., Book of Abstracts, SMRM 1989, Amsterdam, page 23, or in "Human Heart: Tagging with MR Imaging - A method for Noninvasive Assessment of Myocardial Motion", E.A. Zerhouni et al., Radiology, October 1988, pp. 59- 63.

In all these methods it is difficult or not at all possible to evaluate the motion quantitatively, especially automatic evaluation using computer means, due to a relatively poor grid contrast. For images obtained at later stages of the heart cycle the contrast is reduced due to longitudinal relaxation effects. Not only does the grid pattern decay with longitudinal relaxation, but the image intensity becomes increasingly dominated by magnetization which has returned to its relaxed state, and which hence again becomes available for imaging. This relaxed magnetization which is devoid of the grid pattern, and which is thus not useful as far as the tagging is concerned, is a significant factor in reducing the grid contrast.

It is an object of this invention to provide an improved tagging magnetic resonance imaging method which allows for automatic detection of tagging patterns.

A magnetic resonance imaging method according to the present invention is characterized in that the pulse and gradient sequences are applied in pairs with spatially different tagging patterns, images corresponding to the pairs being subtracted so as to form a tagged image.

In such a method according to the present invention, by appropriate choice of the tagging patterns, the component of the image caused by the relaxed magnetization is eliminated and further, as a byproduct, untagged images are obtained by addition of the paired images. The high contrast images obtained allow for automatic grid pattern detection by searching for lines of minima within the tagged image.

In an embodiment of a magnetic resonance imaging method according to the present invention, the tagging patterns in the respective pairs are inverted with respect to each other. Optimum results are obtained with this choice of tagging patterns.

A further embodiment of a magnetic resonance

imaging method according to the present invention is characterized in that the tagging section comprises a first and a second radio frequency pulse separated by a first magnetic field gradient pulse for applying spatial modulation to a component of magnetization perpendicular to the part to be imaged. In this embodiment, the present invention improves the tagging in the cited Axel article, the so-called SPAMM tagging section (a periodic SPACial Modulation of Magnetization prior to imaging), so as to allow for automatic detection of tagging patterns. In this embodiment the SPAMM tagging section achieves a stripe-like tagging pattern.

A further embodiment of a magnetic resonance imaging method according to the present invention is characterized in that the tagging section further comprises a third and a fourth radio frequency pulse separated by a second magnetic field gradient pulse orthogonal to the first magnetic field gradient pulse. A grid-like tagging pattern is obtained in this embodiment.

A further embodiment of a magnetic resonance imaging method according to the present invention is characterized in that in the inverted tagging pattern one of the phases of the radio frequency pulses is in opposition to the phases of the other radio frequency pulses. This achieves optimum mode tagging patterns in SPAMM-like tagging sections.

A further advantageous embodiment of a magnetic resonance imaging method according to the present invention is characterized in that the radio frequency pulses are combined  $180^\circ$  pulses and that the modulus of the image is used for defining the tagging pattern. With this embodiment very sharp minima are obtained, and hence a good motion resolution. Furthermore, the tagging sequence can be kept relatively short, so that the contrast loss due to longitudinal relaxation is negligible.

A further embodiment of a magnetic resonance imaging method according to the present invention is characterized in that the part is a heart, that the tagging section is applied upon detection of the R-wave of the electrocardiogram of the heart, and that the slice selective imaging segments are temporally spaced over a cardiac cycle. The tagging magnetic resonance imaging method according to the present invention is particularly useful for monitoring motion in the heart. With this embodiment multi heart phase images are obtained in which the tagging pattern can be detected automatically by searching for lines of minima within the images. The tagging patterns hardly contain any anatomical contrast. By varying the interval between the R-wave and the application of the tagging section, images of different phases of the heart cycle are

obtained. The later stages of the heart cycle are imaged with respect to the R-wave, the more important a good tagging pattern contrast becomes due to the ever present longitudinal relaxation.

The features and advantages of the magnetic resonance imaging method according to the present invention will be more clearly apparent from the following detailed description, in conjunction with the accompanying drawing, wherein,

Fig. 1 shows, diagrammatically, a magnetic resonance imaging device according to the present invention,

Fig. 2A shows a known multi heart phase myocardial tagging sequence,

Fig. 2B shows a SPAMM tagging section for applying a grid-like tagging pattern.

Figure 1 shows, diagrammatically a magnetic resonance device 1 according to the invention comprising means 2 for generating a stationary homogeneous magnetic field, means 3 for radiating radio frequency pulses to an object 4, modulating means 5 for modulating the rf-pulses, receiving and demodulating means 6 for receiving and demodulating excited magnetic resonance signals comprising sampling means 7 for sampling the magnetic resonance signals, and control means 8 for controlling said means 3, 5, 6 and 7. The device 1 further comprises processing means 9 for processing the sampled magnetic resonance signals, and gradient generating means 10 for generating magnetic field gradients which are superimposed upon the stationary magnetic field. In further detail, the radiating means 3 comprise the modulating means 5, a power amplifier 11, a directional coupler and a transmitter and receiver coil 13. The modulating means 5 comprise a modulator 14 and an oscillator 15. The modulation can be amplitude, frequency or phase modulation or a combination thereof, controlled by the control means 8. The irradiating means 3 excite nuclear spins in the object 4 when transmitting rf-pulses, and the receiving and demodulating means 6 will receive magnetic resonance signals in response thereto. The transmission and reception may have various embodiments, e.g. the coil 13 may be a transmitter/receiver coil or may be embodied as separate transmitter and receiver coils. In the separate embodiment no directional coupler 12 will be present. The receiving and demodulating means 6 may comprise a quadrature demodulator, in which case the sampling means 7 will comprise two analog-to-digital convertors supplying quadrature detected samples of respective sin and cos components of the magnetic resonance signal. The gradient generating means 10 comprise a power supply controlled by the control means 8, for separate control of gradient magnetic coils 17, 18 and 19 which generate magnetic field gradients  $G_x$ ,  $G_y$ ,

and  $G_z$  respectively. The gradient coils 17, 18 and 19 may be arranged such that the field direction of the magnetic field gradients coincides with the direction of the stationary homogeneous magnetic field, and that the directions of the gradients are perpendicular to each other, indicated in figure 1 with mutually perpendicular axes  $x$ ,  $y$  and  $z$ . If the magnetic resonance device 1 is put into operation and the object 4 is placed inside the means 2 for generating the stationary field, a small excess of nuclear spins (of nuclei exhibiting a magnetic moment) will be directed in the direction of the stationary magnetic field, which can be considered macroscopically as a magnetization  $M_0$ , an equilibrium magnetization. The equilibrium magnetization  $M_0$  can be disturbed by the application of pulse and gradient sequences to the object 4. For obtaining physiological information from the object 4, the magnetic resonance device 1 comprises several sensors, such as sensors  $c$ , for picking up cardiac signals which are fed to preamplifiers 20 and 21. The preamplified cardiac signals are fed to an ECG trigger unit 22 via a fibre optic cable 23 by means of an ECG transmitter 24. The ECG trigger unit 22 supplies trigger pulses  $t_r$  to the control means 8 for triggering pulse and gradient sequences. The ECG trigger unit 22 is further controlled by the control unit 8 by means of control signal  $ctl$  for inhibiting the ECG trigger unit 22 from sending pulses during predetermined periods. The processing means 9 may comprise programming means so that the pulse and gradient sequence starts from a predetermined time from the trigger pulse  $t_r$ . The trigger pulse  $t_r$  may be derived in the trigger unit 22 from the R-wave of the electrocardiogram obtained from the object 4. For a more detailed description of a MRI device and a general description of the principles of MRI, reference is made to the handbook, "Practical NMR Imaging", M.A. Foster and J.M.S. Hutchinson, 1987 IRL Press.

In figure 2A, a known multi heart phase myocardial tagging sequence  $tsq$  is shown as a function of the time  $t$ . At  $t = t_0$  a tagging section TAG is applied, e.g. a SPAMM tagging section as described in the cited Axel article. The tagging section is applied under the control of the control unit 8 a predetermined time interval after the control unit receives a trigger pulse  $t_r$  from the trigger unit 22. For obtaining a stripe-like tagging pattern, the tagging section comprises two radio frequency pulses separated by a "wrap" gradient. For obtaining a grid-like tagging pattern, the tagging section may comprise two such rf-pulse gradient combinations in succession. The two one dimensional tagging sections must be separated by a so-called spoiler or dephasing gradient, to obtain a multiplication of these two tagging patterns. The tagging section TAG is followed by a multi heart

phase imaging section after a time interval  $\delta t$ , the imaging section comprising a number of gradient echo imaging segments temporally spaced over the cardiac cycle. Each imaging segment comprises an rf-pulse P1 with excitation angle  $\alpha$ , a slice selection gradient G1, a phase encoding gradient G2 and a readout gradient G3 applied by the rf-coil 13 and the gradient coils 17, 18 and 19, respectively. The pulses  $\alpha_k$ ,  $k=0, \dots, n-1$  ( $n$ : total number of heart phase images), are low flip angle pulses, not necessarily all equal. The respective pulse angles can be optimized for particular repetition times and tissue longitudinal relaxation times. The sequence  $tsq$  is repeated a number of times, e.g. 256, for different values of the phase encoding gradient G2. The received and sampled magnetic resonance signals around  $t = t_k$ ,  $t = t_{k+1}$ , ... are processed by the processing means 9, a tagged image being obtained for each heart phase. The time interval between the  $k$ th and  $(k+1)$ th imaging segment is  $\delta t_k$ .

In figure 2B, a SPAMM tagging section TAG for applying a grid like tagging pattern is shown, comprising first and second radio frequency pulses P2 and P3 separated by a first "wrap" gradient G7, and comprising third and fourth radio frequency pulses P4 and P5 separated by a second "wrap" gradient G6, the first and the second "wrap" gradient being orthogonal with respect to each other. The gradients G4 and G5, which must be in slice selection direction, are applied to dephase any  $x$  and  $y$  components of the tagging magnetization.

The spatial modulation can be described as

$$TAG(x,y) = \cos(\beta x) \cdot \cos(\gamma y)$$

The factors  $\beta$  and  $\gamma$  are determined by the integrals of the gradients G7 and G6, respectively. The amplitudes of the pulses P2, P3, P4 and P5 can be constant. The radio frequency pulses P2 and P3, and P4 and P5 form so-called "11" sets of pulses then, the sum of the pulse angles of the pulses P2 and P3, and P3 and P4 being in a range of  $90^\circ$  and  $180^\circ$ . Binomial pulse combinations such as "121" or "1331" pulses can also be applied, separated by gradients. Practically the summed angle amounts between  $90^\circ$  and  $180^\circ$ . For a specified tissue, a specified number of excitation pulses of a specified angle and for a specified time point, the tagging angle can be optimized to obtain a better contrast than a known SPAMM sequence. For example, when measuring a diastolic, a midstolic, and a systolic modulus image with an excitation angle of  $35^\circ$ , the tagging contrast at the systolic images on the myocardium is maximal, using a tagging angle of about  $140^\circ$ . The present invention overcomes such problems. Application of binomial pulse combinations might take

too long, i.e. the tagging section might not be completed before the heart begins to move.

Before a more detailed description of the present invention will be given, shortcomings of the known SPAMM tagging will first be outlined. It is assumed for simplicity that the object 4 to be imaged has a uniform magnetization density equal to unity. Then, immediately after the TAG section, the z-magnetization is given by

$$M_0 \cdot \text{TAG}(x,y)$$

The shape of each heart phase image is determined by the z magnetization immediately before the respective rf excitation pulses P1. For the first image this z-magnetization is

$$1 - (1 - M_0 \cdot \text{TAG}(x,y)) \cdot \exp(-\delta T/T_1) = M_0 \cdot \text{TAG}(x,y) \cdot \exp(-\delta T/T_1) + (1 - \exp(-\delta T/T_1))$$

where  $T_1$  represents the longitudinal relaxation time.

In the above formula, the first term

$$QT_0 = M_0 \cdot \text{TAG}(x,y) \cdot \exp(-\delta T/T_1)$$

represents a decaying image of the object which has been modulated by the tagging pattern, and the second term,

$$QR_0 = 1 - \exp(-\delta T/T_1)$$

represents an image of the object due only to that magnetization that has been relaxed since the application of the tagging section TAG. For the kth heart phase image

$$QT_k = QT_0 \cdot \prod_j \cos(\alpha_j) \cdot \exp(-\delta_j/T_1), \text{ for } j=0 \text{ to } k-1$$

and, recursively,

$$QR_{k+1} = 1 - (1 - QR_k) \cdot \exp(-\delta k/T_1)$$

$QT_k$  contains the information for monitoring myocardial motion, while  $QR_k$  is a term which reduces the contrast of the tagging grid, and for a tagging section with a  $90^\circ$  tagging angle it can easily be shown that

$$QT_k = \lambda k \cdot (1 - \cos(\beta x)) \cdot (1 - \cos(\gamma y))$$

$\lambda k$  being dependent on the  $\alpha_j$ 's,  $\delta_j$ 's,  $T_1$  etc., but not depending on the spatial coordinates  $x$  and  $y$ . In the presence of motion the variables  $x$  and  $y$  represent object coordinates at the point in time  $t=t_0$  at which the tagging section TAG is applied. The tagged image is given by

$$\lambda k \cdot (1 - \cos(\beta x)) \cdot (1 - \cos(\gamma y)) + QR_k$$

Basically the tagging grid can be automatically detected by searching for the lines of minima within the tagged image. However due to the presence of the term  $QR_k$  and the influence of  $\lambda k$ , the actual values of these minima vary widely. In an image with large intensity variations, which is usually the case, it could be very difficult to detect these minima.

For a tagging section with a  $180^\circ$  tagging angle,

$$QT_k = \lambda k \cdot \cos(\beta x) \cdot \cos(\gamma y)$$

And for modulus type images, as is dealt with in many cases, the image shape is as follows,

$$|\lambda k \cdot \cos(\beta x) \cdot \cos(\gamma y) + QR_k|$$

In the latter case it does not suffice to search for lines of minima, their position being influenced by  $\lambda k$  and  $QR_k$ . The tagging grid, which is positioned between the lines of minima, can only be assigned definitely in homogeneous regions of the object in this case.

According to the present invention, the term  $QR_k$  is eliminated in order to allow for reliable automatic detection of tagging grids in tagged images. The elimination is achieved by performing pairs of measurements with two spatially differing tagging functions and by subtracting the tagged images resulting from two tagging pulse and gradient sequences  $tsq$  with different tagging sections TAG incorporating these tagging functions. Denoting the respective tagging functions by  $TAG1(x,y)$  and  $TAG2(x,y)$ , the best results are obtained when

$$TAG1(x,y) = -TAG2(x,y)$$

For a  $180^\circ$  SPAMM tagging section TAG as shown in figure 2B

$$TAG1(x,y) = \cos(\beta x) \cdot \cos(\gamma y)$$

This tagging pattern can be inverted by negating the phase of any one of the four radio frequency pulses P2, P3, P4 or P5, i.e. by replacing one of the four  $90^\circ$  pulses in "11" sets of pulses P2, P3 and P4, P5 respectively by a  $-90^\circ$  pulse. With this choice of tagging functions, the form of the tagged image is as follows

$$|2\lambda k \cdot \cos(\beta x) \cdot \cos(\gamma y)|$$

By addition of the two tagged images resulting from the two tagging sequences an image with  $T_1$  dependent anatomical contrast will be obtained with the form of

2.QR<sub>k</sub>

According to the present invention, the lines of minima substantially have zero intensity, which make them very easy to detect. The zero intensity lines are detected automatically by means of the processing means 9 through scanning of the resulting tagged image. Furthermore, the tagging magnetic resonance method according to the present invention gives very sharp minima when the tagging grid is defined as a modulus image.

In case of 90° tagging angles, according to the present invention, the tagging grid lines can be narrowed by applying binomial pulses, with a summed pulse angle of 90° for each tagging direction. The form of the tagging function then is, by rough approximation

$$(1 - \cos^n(\beta x)) \cdot (1 - \cos^n(\gamma y))$$

n being the order of the binomial sequence. The duration of the tagging sequence increases with the order n and tagging might not be completed before the heart starts to move. In case of myocardial motion the 180° variant is thus preferred.

## Claims

1. A magnetic resonance imaging method for monitoring motion of a part of an object (4) to be imaged by application of pulse and gradient sequences (tsq) to the object (4), which sequences (tsq) comprise a tagging section (TAG) for producing a tagging pattern and an imaging section (P1, G1, G2, G3) for producing magnetic resonance signals in the part of the object, the resonance signals being used for forming at least one image of the object, characterized in that the pulse and gradient sequences (tsq) are applied in pairs with spatially different tagging patterns, images corresponding to the pairs being subtracted so as to form a tagged image.
2. A magnetic resonance imaging method as claimed in Claim 1, characterized in that the tagging patterns in the respective pairs are inverted with respect to each other.
3. A magnetic resonance imaging method as claimed in Claim 1 or 2, characterized in that the tagging section comprises a first (P2) and a second (P3) radio frequency pulse separated by a first magnetic field gradient pulse (G7) for applying spatial modulation to a component of magnetization perpendicular to the part to be imaged.
4. A magnetic resonance imaging method as claimed in Claim 1, 2 or 3, characterized in that the tagging section (TAG) further comprises a third (P4) and a fourth (P5) radio frequency pulse separated by a second magnetic field gradient pulse (G6) orthogonal to the first magnetic field gradient pulse (G7).
5. A magnetic resonance imaging method as claimed in Claim 3 or claim 4, characterized in that in the inverted tagging pattern one of the phases of the radio frequency pulses (P2, P3, P4, P5) is in opposition to the phases of the other radio frequency pulses.
6. A magnetic resonance imaging method as claimed in any one of the Claims 3 to 5, characterized in that the radio frequency pulses (P2, P3, P4, P5) are combined 180° pulses (P2 & P3, P4 & P5) and that the modulus of the image is used for defining the tagging pattern.
7. A magnetic resonance imaging method as claimed in any one of the preceding Claims, characterized in that the tagging patterns are detected by searching for lines of minima within the image.
8. A magnetic resonance imaging method as claimed in any one of the preceding Claims, characterized in that the imaging section (TAG) comprises at least one slice selective imaging segment (P1, G1, G2, G3) for obtaining at least one slice image of the part.
9. A magnetic resonance imaging method as claimed in Claim 8, characterized in that the part is an heart, that the tagging section is applied upon detection of the R-wave of the electrocardiogram of the heart, and that the slice selective imaging segments are temporally spaced over a cardiac cycle.
10. A magnetic resonance imaging device (1) for monitoring motion of a part of an object (4), which imaging device (1) comprises means (2) for generating a stationary homogeneous magnetic field, means (3) for radiating radio frequency pulses to the object (4) which is placed in the stationary field, gradient generating means (10) for generating magnetic gradient fields which are superimposed upon the stationary field, and processing means (9) which are arranged for applying pulse and gradient sequences (tsq) to the object (4) under the control of control means (8); the sequences comprising a tagging section (TAG) and an

imaging section (P1, G1, G2, G3), and which imaging device (1) further comprises receiving means (6) and sampling means (7) for receiving and sampling of magnetic resonance signals generated by the sequences (tsq) in at least a part of the object (4), the processing means (9) being arranged for processing sampled resonance signals into at least one image, characterized in that the processing means (9) are further arranged for applying the pulse and gradient sequences (tsq) in pairs with spatially differing tagging patterns, for subtracting images corresponding to the pairs so as to form a tagged image, and for monitoring the motion in the part by scanning the tagging pattern in the tagged image.

5

10

15

20

25

30

35

40

45

50

55

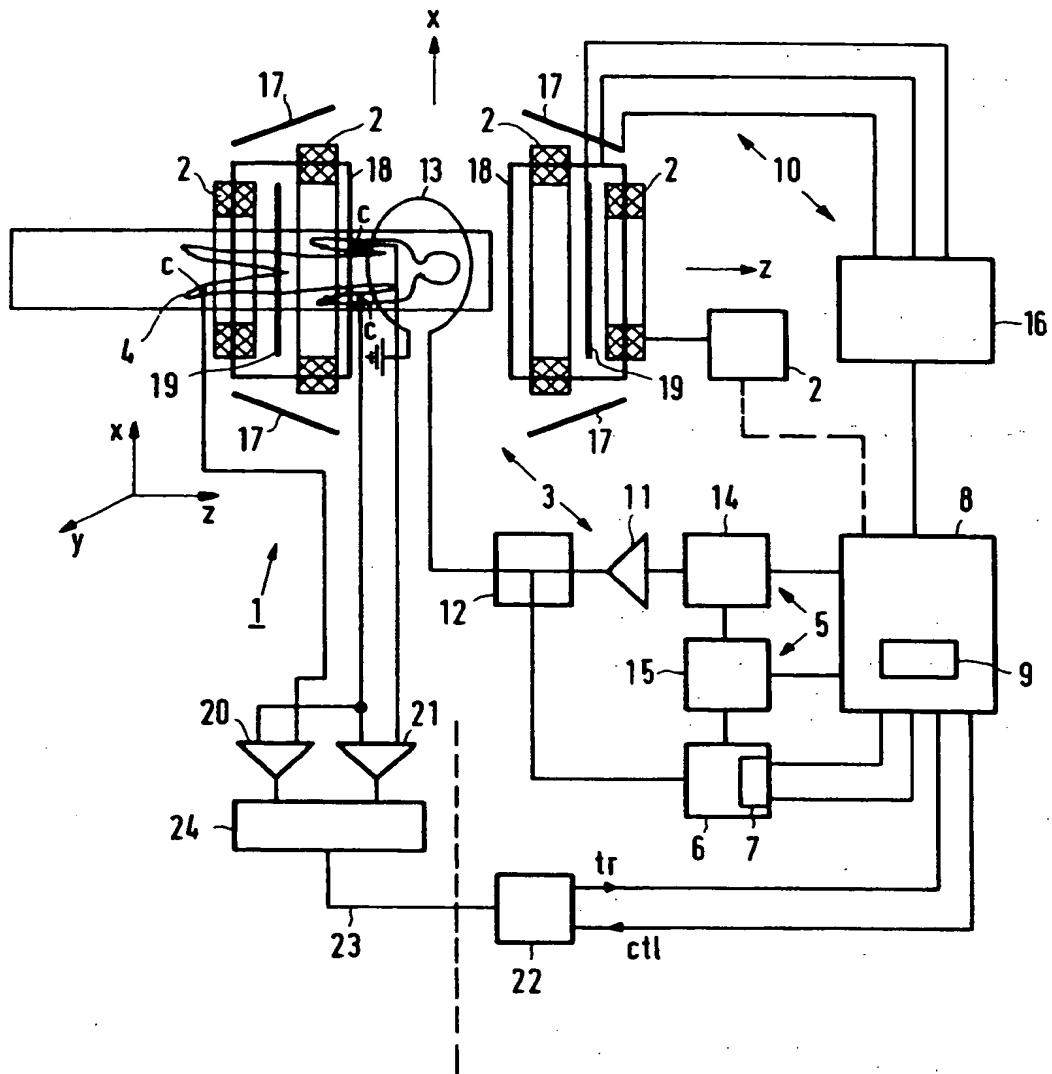


FIG.1



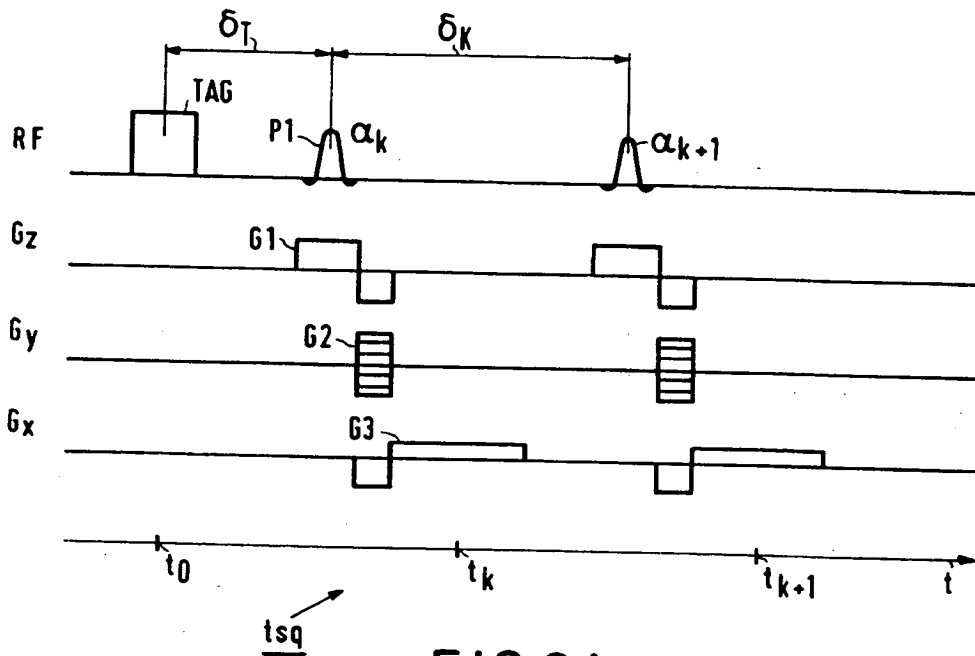


FIG. 2A

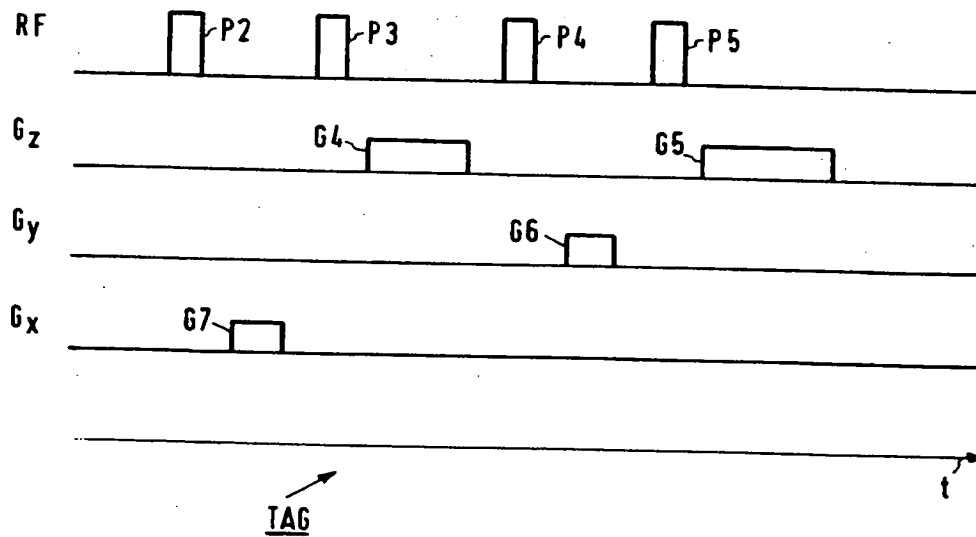


FIG. 2B

FOR NOT GO VICTIMS

*Handwritten signature*

(19)



Europäisches Patentamt  
European Patent Office  
Office européen des brevets



(11) Publication number:

**0 507 392 A3**

(12)

**EUROPEAN PATENT APPLICATION**

(21) Application number: 92200885.9

(51) Int. Cl.<sup>5</sup> **G01R 33/48, G01R 33/56**

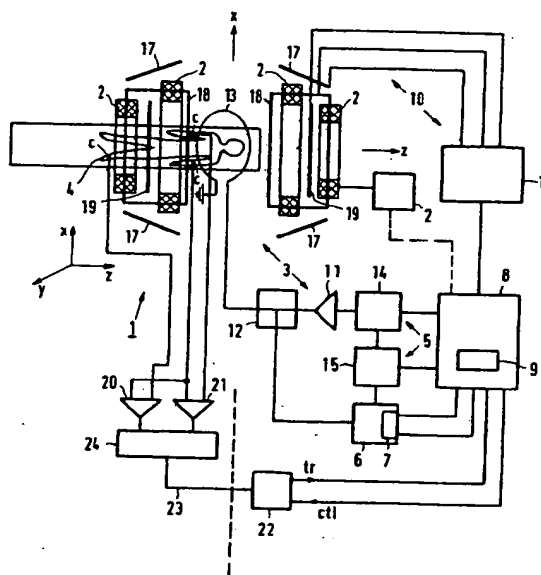
(22) Date of filing: 30.03.92

(30) Priority: 02.04.91 EP 91200762

(43) Date of publication of application:  
07.10.92 Bulletin 92/41(94) Designated Contracting States:  
**DE FR GB NL**(68) Date of deferred publication of the search report:  
23.06.93 Bulletin 93/25(71) Applicant: **N.V. Philips' Gloeilampenfabrieken**  
**Groenewoudseweg 1**  
**NL-5621 BA Eindhoven(NL)**(72) Inventor: **KcKinnon, Graeme**  
**c/o INT. OCTROOIBUREAU B.V. Prof.****Holstlaan 6****NL-5656 AA Eindhoven(NL)**Inventor: **Fischer, Stefan****c/o INT. OCTROOIBUREAU B.V. Prof.**  
**Holstlaan 6****NL-5656 AA Eindhoven(NL)**Inventor: **Boesiger, Peter****c/o INT. OCTROOIBUREAU B.V. Prof.**  
**Holstlaan 6****NL-5656 AA Eindhoven(NL)**(74) Representative: **De Jongh, Cornelis**  
**Dominicus et al**  
**INTERNATIONAAL OCTROOIBUREAU B.V.**  
**Prof. Holstlaan 6**  
**NL-5656 AA Eindhoven (NL)**(54) **Magnetic resonance imaging method and device for monitoring motion of a part of an object.**

(57) Magnetic resonance imaging methods for myocardial tagging are known for assessment of myocardial motion. In such methods tagging pulse and gradient sequences (tsq) are applied to the heart of an object (4) for acquiring tagged magnetic resonance images of, for example, heart slices from diastole to systole, the sequences being triggered upon detection of the R-wave of an ECG. With these known sequences, such as the so-called SPAMM sequence, it is very difficult, if at all possible, to automatically detect the tagging grid and therefore to evaluate the motion quantitatively.

According to the present invention, a magnetic resonance imaging method and device (1) are disclosed which allow for automatic quantitative motion evaluation of a part of an object (4). The pulse and gradient sequences (tsq) are applied in pairs with spatially differing tagging patterns, images corresponding to the pairs being subtracted so as to form a tagged image. The motion is monitored by scanning of the tagging pattern in the tagged image.

**FIG. 1****EP 0 507 392 A3**



European Patent  
Office

# EUROPEAN SEARCH REPORT

Application Number

EP 92 20 0885  
Page 1

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
A	MAGNETIC RESONANCE IN MEDICINE. vol. 15, no. 2, 1 August 1990, DULUTH, MN US pages 334 - 339 T.J. MOSHER ET AL. 'A DANTE Tagging Sequence for the Evaluation of Translational Sample Motion' * the whole document *	1,3,8,10	G01R33/48 G01R33/56
A,D	RADIOLOGY vol. 169, no. 1, 1 October 1988, NEW YORK, US pages 59 - 63 E.A. ZERHOUNI ET AL. 'Human Heart: Tagging with MR Imaging - A Method for Noninvasive Assessment of Myocardial Motion' * the whole document *	1,10	
A,D	RADIOLOGY vol. 171, no. 3, 1 June 1989, NEW YORK, US pages 841 - 845 L. AXEL ET AL. 'MR Imaging of Motion with Spatial Modulation of Magnetization' * the whole document *	1,3,10	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			G01R
A	US-A-4 953 554 (E. ZERHOUNI ET AL.) * column 1, line 25 - column 2, line 2 * * column 6, line 3 - column 7, line 25 *	1,8-10	
A	EP-A-0 205 136 (GENERAL ELECTRIC COMPANY) * page 7, line 13 - page 13, line 22 *	1,8-10	
A	DE-A-3 918 625 (HITACHI, LTD.) * abstract * * column 5, line 35 - column 6, line 42 *	1,8-10	
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 26 APRIL 1993	Examiner HORAK G.I.
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons A : member of the same patent family, corresponding document	

EPO FORM 1503 01.82 (P0601)



European Patent  
Office

# EUROPEAN SEARCH REPORT

Application Number

EP 92 20 0885  
Page 2

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
A,P	WO-A-9 203 089 (THE TRUSTEES OF THE UNIVERSITY OF PENNSYLVANIA) * page 8, line 1 - page 13, line 14 * * page 16, line 10 - page 19, line 35 * ---	1,3,8-10	
A,P	US-A-5 054 489 (L. AXEL ET AL.) * column 1, line 7 - line 68 * * column 2, line 38 - column 3, line 37 * -----	1,3,8-10	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 26 APRIL 1993	Examiner HORAK G.I.
<b>CATEGORY OF CITED DOCUMENTS</b> X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons ----- & : member of the same patent family, corresponding document			

EPO FORM 1503 (11.92) (P0601)

**THIS PAGE BLANK (USPTO)**